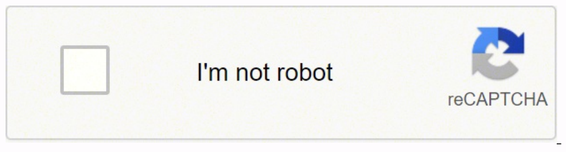

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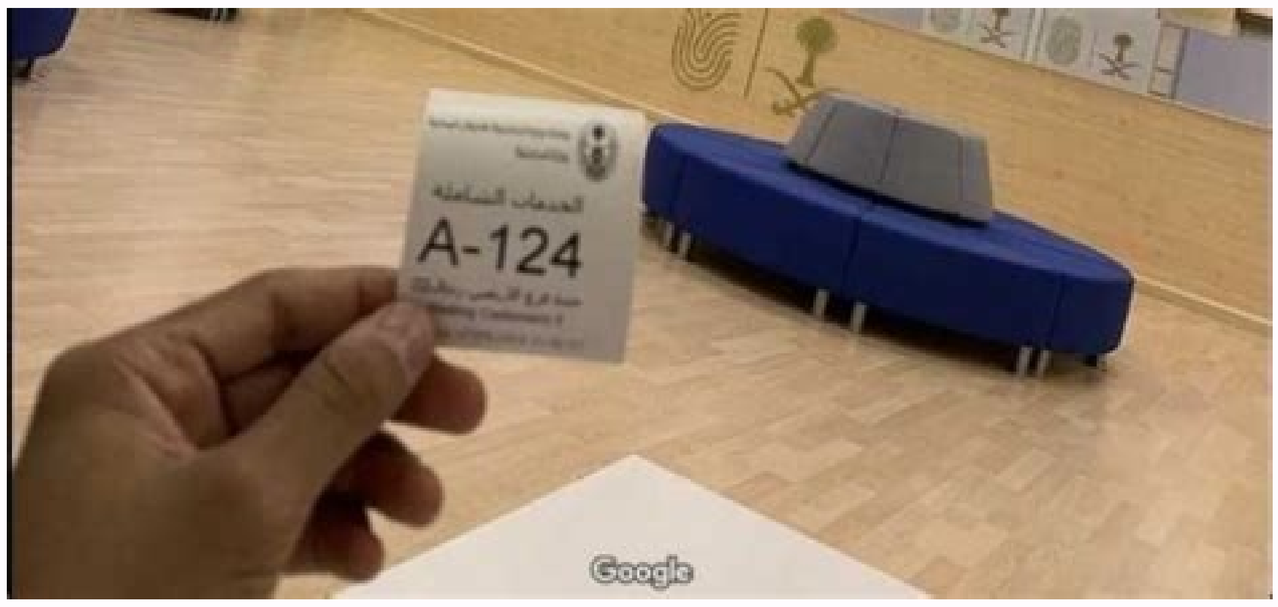

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AFAIRS or AHWAL cellular certification procedure The procedure for obtaining a birth certificate from Babybird in Saudi Arabia is quite easy, just a job from one to two hours. However before I go, you must be ready with all the necessary documents. Trends: Experience in the Birth Certificate shared by Riyadh Chico Compulsory documents to obtain Certificate of Birth of Civil Affairs (Ahwal al Madni): 1. Appointment of Ahwal Al Madni or Civil Affairs 2. Original and copy of the Hospital Birth Notification o Ministry of Health, you can get this from the hospital where your baby was born. 3. Original and copies of IQamas of father and mother of announcement. 4. Original and passport copies of father and mother. 5. Civil Affairs Birth Certificate Form, you can fill it with the help of anyone who knows Arab. Procedure to obtain the Civil Affairs Birth Certificate (AHWAL AL Madni): - Reach the Ahwal Office Al Madni one hour before your date of date, show the documents at the entrance and get the first token, at the counter that It will organize your documents then it will give you second token. Try writing your baby name in English in Hospital Certificate to avoid spelling errors. Similar: Arrival visas for just born at Saudã Airport - In the second counter you will take all the documents and will issue a birth certificate for your baby, it is green in color. - Your birth certificate will have details of your baby-born baby including the iqama number and the name in English. - Take the third tab for the MOFA certificate that is at the other counter, it will charge 30 SR for MOFA seal. See also: the employer's duty to provide medical insurance for the employee and family of it-now you can see your baby name and the Iqama number of baby in Absher, her name of the baby the father’s name on it. - You can now apply for your baby’s passport at your Embassy in Saudi Arabia. Note: You must complete the above procedure within 1 month of your baby’s birth, If you you One month you have to pay a fine of 50 SR to Civil Affairs (Ahwal Al Madni). Six years ago, a teenager from Italy traveled to the United States in the hope of finding a diagnosis for her mysterious disease, which had made her unable to walk and forced her to have a respiratory tube. Now, researchers have diagnosed adolescent Claudia Digregorio and 10 other children with a new form of amyotrophic lateral sclerosis (ALS) that affects children and progresses slower than usually seen with this condition.In addition, researchers have identified a gene that appears to cause this form of ALS, and may also have identified a possible treatment for the condition. “We hope these results will help doctors recognize this new form of ALS and lead to the development of treatments that will improve the lives of these children and young adults”, says Dr. Carsten BÃ¶nnemann, senior researcher at the National Institute of Neurological Disorders and Stroke (NINDS). and a lead author of the study, he said in a statement. “We also hope that our results may provide new clues to understand and treat other forms of the disease”.Related: 10 things you didn’t know about the brainALS is a rare disease that causes progressive degeneration and death of nerve cells that control voluntary muscle movements, such as chewing. walking, speaking and breathing, according to the NINDS. Most people with ALS develop symptoms between the ages of 55 and 75, and the disease usually progresses rapidly, with death occurring between three and five years after diagnosis.But with the new form of ALS, the symptoms appeared much earlier, often around 4 years of age. For many of the 11 patients, the first signs of the disease were trouble walking and spasticity in the lower extremities. In the Many of the patients, such as Digregorio, needed a wheelchair to move around. move. a tracheotomy tube to breathe support. before the digregory went to the uu, the then 15 years of age met with the French pope, who offered prayers for his health, according to a question of 2015 of the Nih registration, an information bulletin of the national health institutes. the researchers found an answer. for digregory, which was the first patient included in the study. Despite development symptoms at such an early age, she and the other 10 children showed signs of als flash in neurological exams, including severely weakened or paralyzed muscles, researchers said. "These young patients had many of the problems of superior and lower motor neuron that are indicative of als," said Dr. payam mohassel, a member of the clinical research in the nih. (motor neurons are nerve cells in the brain and spinal cord that send signals that control the movement. superior motor neurons originate in the brain and send signals to the lower engine neurons, which are in the spinal cord.) "What made these cases unique was the early start-up age and slower progression of symptoms," said mohassel. "this asked us to wonder what this different form of als was underlying."With genetic sequencing, researchers found that these patients had genetic changes in a specific section of a gene called spltc1. This gene is involved in the production of fats called spurgolipids, which are particularly abundant in the brain tissue. a series of other deadly neurodegenerative diseases, including nieman selection diseases and Tay-Sachs disease, are also caused by problems with the metabolism of sfingolypse. more research revealed that the mutation in spltc1 increased the levels of sphingolipids. specifically, researchersthat the mutation "reducing the brake" of an enzyme involved in the production of sphingolipids, which means that the body continues to shake these fats without its usual feedback system to tell you when it stops. The findings mean that the restoration of this "Frake" could treat this type of ALS, the authors said. In anotherresearchers tested a therapy called small interfering RNA, or siRNA, in which small strands of RNA work to “silence” a mutated gene, in this case, SPLTC1. In studies conducted on laboratory dishes, the therapy worked to restore sphingolipid levels to normality”.Our ultimate goal is to translate these ideas into effective treatments for our patients who currently don’t have therapeutic options”, BÃ¶nnemann said.Future studies should also investigate whether problems with sphingolipid metabolism play a role in other forms of ALS, the authors said. Live science. Science.Ã Filled form 87 for Birth Certificate. A copy of the Saudi visa stamped on the mother’s passport (in case the mother does not have an Iqama). A screenshot of the wife’s details from Absher (in case the wife is on a family visit visa). Visit Ahwal Madani Second Floor, North Tower Jawad Center, on top of Ahwal Madani Corner King Fahad Road and Damar Bin Taalabah Street 21491 Jeddah Saudi Arabia ...

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